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Practitioner's Docket No. MBIO1997-018DV1ACN1(M) *PATENT* 0360

BX SEQ

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of: MBIO1997-018DV1ACN1(M)
Application No.: 09/993,179 Group No.: 1636
Filed: November 6, 2001 Examiner:
For: SECRETED PROTEINS AND NUCLEIC ACIDS ENCODING THEM
Inventors: Sean A. McCarthy et al.

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SUBMISSION OF "SEQUENCE LISTING," COMPUTER READABLE COPY, AND/OR AMENDMENT PERTAINING THERETO FOR BIOTECHNOLOGY INVENTION CONTAINING NUCLEOTIDE AND/OR AMINO ACID SEQUENCE

Certificate of First Class Mailing (37 CFR 1.8(a))
I hereby certify that this correspondence is being deposited with the United States Postal Service as first class mail in an envelope addressed to: Assistant Commissioner for Patents, Washington, D.C. 20231 on the date set forth below.

January 30, 2002 By: Jean Hunziker
Date of Signature and of Mail Deposit Sean Hunziker

1. This replies to the Office Communication including a Notice to Comply with Requirements for Patent Applications Containing Nucleotide and/or Amino Acid Sequence Disclosures dated January 3, 2002. This response corrects the errors noted in the Raw Sequence Listing Error Report which accompanied the Notice to Comply with Requirements for Patent Applications Containing Nucleotide Sequence and/or Amino Acid Sequence Disclosures.

   A copy of the Notice to Comply with Requirements for patent Applications Containing Nucleotide and/or Amino Acid Sequence Disclosures and a copy of the Raw Sequence Listing Error Report is enclosed.

IDENTIFICATION OF PERSON MAKING STATEMENT

2. I, Jean M. Silveri state the following:

ITEMS BEING SUBMITTED

3. Submitted herewith are:

Listing(s)" for the nucleotide and/or amino acid sequence(s) in this application. Each equence Listing" is assigned a separate identifier as required in 37 C.F.R. Section 821(c) and 37 C.F.R. Sections 1.822 and 1.823.

each "Sequence Listing" submitted for this application in computer readable form, in cordance with the requirements of 37 C.F.R. Sections 1.821(e) and 1.824.

nt that the content of each "Sequence Listing" submitted and each computer readable re the same, as required in 37 C.F.R. Section 1.821(g).

his submission is made in fulfilling the requirement under 37 C.F.R. Section 1.821(g), ment that the submission includes no new matter.

STATEMENT THAT "SEQUENCE LISTING" AND COMPUTER READABLE COPY ARE THE SAME OR THAT PAPERS SUBMITTED INCLUDES NO NEW MATTER

uter readable form submitted in this application, including those forms requested to be rred from applicant's other application, is the same as the "Sequence Listing" to which it ated to relate.

ccompanying this submission, or for which a request for transfer from applicants' other tion, introduce no new matter.

STATUS

er than a small entity.

MILLENNIUM PHARMACEUTCALS, INC.

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SIGNATURE OF PRACTITIONER

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COPY

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APPLICATION NUMBER	FILING/RECEIPT DATE	FIRST NAMED APPLICANT	ATTORNEY DOCKET NUMBER
09/993,179	11/06/2001	Sean A. McCarthy	MBIO1997-018DV1ACN1 (M)

CONFIRMATION NO. 1059

FORMALITIES LETTER

OC000000007254847

Millennium Pharmaceuticals, Inc.
75 Sidney Street
Cambridge, MA 02139

Date Mailed: 01/03/2002

NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

Applicant is given **TWO MONTHS FROM THE DATE OF THIS NOTICE** within which to file the items indicated below to avoid abandonment. Extensions of time may be obtained under the provisions of 37 CFR 1.136(a).

- A copy of the "Sequence Listing" in computer readable form has been submitted. However, the content of the computer readable form does not comply with the requirements of 37 C.F.R. 1.822 and/or 1.823, as indicated on the attached copy of the marked -up "Raw Sequence Listing." Applicant must provide a substitute computer readable form (CRF) copy of the "Sequence Listing" and a statement that the content of the sequence listing information recorded in computer readable form is identical to the written (on paper or compact disc) sequence listing and, where applicable, includes no new matter, as required by 37 CFR 1.821(e), 1.821(f), 1.821(g), 1.825(b), or 1.825(d).

For questions regarding compliance to these requirements, please contact:

- **For Rules Interpretation, call (703) 308-4216**
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A copy of this notice MUST be returned with the reply.

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